

Product datasheet for TP726793

OriGene Technologies, Inc.

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Junctional Adhesion Molecule 2 (JAM2) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant Human Junctional Adhesion Molecule B/JAM-B/CD322 (C-6His)

Species: Human

Expression cDNA Clone

or AA Sequence:

Phe29-Asn236

Tag: C-His

Buffer: Lyophilized from a 0.2 um filtered solution of 20mM Tris-HCl,150mM NaCl,pH8.0.

Note: Recombinant Human Junctional Adhesion Molecule B is produced by our Mammalian

expression system and the target gene encoding Phe29-Asn236 is expressed with a 6His tag

at the C-terminus.

Storage: Lyophilized protein should be stored at < -20°C, though stable at room temperature for 3

weeks. Reconstituted protein solution can be stored at 4-7°C for 2-7 days. Aliquots of

reconstituted samples are stable at < -20°C for 3 months.

Stability: 12 months from date of despatch

Locus ID: 58494 **UniProt ID:** P57087

Synonyms: Junctional Adhesion Molecule B; JAM-B; Junctional Adhesion Molecule 2; JAM-2; Vascular

Endothelial Junction-Associated Molecule; VE-JAM; CD322; JAM2; C21orf43; VEJAM

Summary: Junctional Adhesion Molecule B (JAM-B) is a single-pass type I membrane protein that

belongs to the juctional adhesion molecules family. JAM-B includes a signal sequence (aa 1-28), an extracellular region (aa 29-238) with one lg-like C2-type domain and one lg-like V-type domain, a transmembrane segment (aa 239-259), and a cytoplasmic domain (aa 260 - 298). JAMB is localized to the tight junctions between endothelial cells or epithelial cells. JAM-B is prominently expressed in the heart, placenta, lung, foreskin and lymph node. It is also present on the endothelia of other vessels. JAM-B acts as an adhesive ligand for interacting with a variety of immune cell types and may play a role in lymphocyte homing to secondary

lymphoid organs.

Protein Families: Druggable Genome, Transmembrane







Protein Pathways:

Cell adhesion molecules (CAMs), Epithelial cell signaling in Helicobacter pylori infection, Leukocyte transendothelial migration, Tight junction